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Development of a new automatic **on-site** detector of pesticides in natural waters by photo induced fluorescence, application to three phenylurea and benzoylurea herbicide.

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ABSTRACT

Prototypes of on-site automatic photo induced fluorescence detectors of pesticide in natural waters are set up and applied for the determination of the benzoyl- and phenylurea pesticides, namely fluometuron, monolinuron and diflubenzuron. As these pesticides present no native fluorescence the set up system use the photo conversion under UV irradiation of these pesticides into highly fluorescent photoproducts.

A first system, called AUTOPIF, (evolution the commercial AQUAPOD system) is develop using a detection via a diode array spectrometer. To improve the sensitivity of the method, a second system, called AUTOPIF+, is developed with a more resolute spectrometer and an intensified CCD camera detection.

Analytical applications were carried out in aqueous solution and detected on line with the AUTOPIF and AUTOPIF + system. The calibration curves are linear over one order of magnitude, and the limits of detection are in the $\mu\text{g mL}^{-1}$ range. The analytical performances of these methods for the determination of the three pesticides are satisfactory in comparison to other classical PIF methods published for the determination of phenylurea pesticides in aqueous solutions. Our results show that the AUTOPIF and AUTOPIF+ methods are versatile, sensible and can be easily applied as an alert system to detect pollutant residues in natural waters over a threshold value.

1.Introduction

The importance of pesticides in increasing agricultural yields and crops protection is no longer to be demonstrate. However, they can induce accidental pollution of natural waters, disrupt aquatic life of many species and also human health due to their persistence and their toxicity. Indeed, many of these pesticides are water soluble and their soil-based residues can remain for several months following application, they can reach foodstuffs and also ground waters^[1,2]. It becomes so very difficult to escape to the consumption of pesticides. They were real public health problem because cited as responsible to many pathologies such as cancers, neurological, respiratory and reproductive disorders^[3,4,5].

Moreover, the presence of pesticides in ground water will disrupt water treatment plants for drinking water production by contaminating the entire facility and will be difficult to remove from tap water. Consequently, these plants have to be protected permanently by an automatic **on-site** system from an accidental pollution to detect any concentration over and alert threshold determined by the plant's processing capabilities.

As some pesticides present great photochemical reactivity under UV irradiation, they can be transformed into photoproducts which are detected by fluorescence. It corresponds to the Photo Induced Fluorescent (PIF) methods, which have been developed by numerous authors for the analysis of herbicides ^[6,7,8], insecticides ^[9,10,11,12] and others pesticides ^[13,14,15,16,17]. It is worthwhile to note that Diaw et al. ^[18,19] have determinate fenuron and diflubenzuron, by classical-PIF method using UV irradiation and DL-PIF method with laser irradiation, in Senegalese natural water with satisfactory mean recovery percentage values in the range 95-105%. And, on the other hand, in her previous work, Mbaye et al. applied an **on-site** automatic detector (namely "AQUAPOD") for the determination of hydrocarbons in aqueous medium by fluorescence detection to monitor pollution of natural waters ^[20].

In this present work, we have associated these two technics to implement, for the first time, the PIF method on an **automatic on-site detector prototype** (namely "AUTOPIF") for the continuous monitoring of pesticides in natural waters. Two configurations of the system are studied and compared. Both are using a UV lamp to obtain the photoproducts. The first **prototype** analyzes their fluorescence on a diode array spectrometer (AUTOPIF); The second **prototype** use a more resolutive spectrometer and a detection by an ICCD camera (AUTOPIF+) to increase selectivity and sensitivity.

These new systems offer several benefits. Firstly, the PIF method allows to detect non-native fluorescent pesticides. Secondly, using fluorescence measurements instead of absorbance increases the sensitivity and allows a direct of the detection of the pesticides. Thirdly, it avoid the use of complex fluidic system to concentrate the pesticides by solid phase extraction followed by UV detection ("Aquapod SPE50" system ^[21]), or the use of HPLC online separation ("SAMOS" system ^[22]).

Here, we investigated benzoyl- and phenylureas (BPUs) pesticides due to their intensive application in agriculture, such as systemic herbicides photosynthesis inhibitors fluometuron (FLM) and monolinuron (MLN) ^[23,24]; and diflubenzuron (DFB) a non-systemic insect growth regulator ^[25]. The two systems are then used for the determination of the BPUs pesticides (FLM, MLN and DFB) in natural waters. Analytical performances of these methods were studied and also the analytical applications were carried out in tap and natural waters.

2. Material and methods

2.1 Reagents

Technical-grade Monolinuron, Fluometuron and Diflubenzuron (Table 1) (purity > 99%) and spectroscopy solvent Methanol were obtained from Sigma Aldrich (analytical reagent grade). Ultra-pure water (Millipore Mro-MQ System) was used as working solutions.

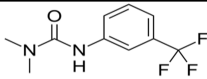
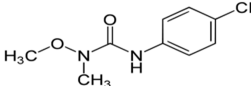
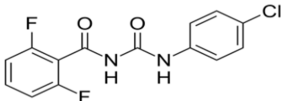
2.2 Materials

Peristaltic pump (Minipuls), SPE C18 cartridge (LiChrolutRP-18E200mg40–63 mm). 30 mL quartz flow tubes (1cm diameter). 0.5 mL cylindrical quartz flow cell. Diode array spectrometer (200-400 nm, Oceans Optics). SpectraPro-550i spectrometer, $f=500\text{mm}$ (Acton, MA, USA). CCD intensified Camera (Princeton instruments, NJ, USA). 125 W Hg lamp (Philips). 280 nm UV LED (Thorlabs, France). Commercial “AQUAPOD LIGHT” system (HOCER, instrumentation company, Brest France). Stagraphics 18 Centurion software.

2.3 Methods

Stock standard solutions of the pesticides (310 mg/L) were prepared by dissolving the compounds in methanol. Serial dilutions were performed to obtain the working standard solutions in water. To obtain natural water matrix samples free of organic contamination, traces of organic compounds are eliminated by preparative chromatography using SPE C18 cartridge. The cartridge was preconditioned with 5mL of methanol followed by 5mL of ultrapure water; 50mL of water samples were passed through the cartridge before being spiked.

Table1 : Chemical properties of fluometuron, Monolinuron and diflubenzuron

Pesticides	Chemical structure	Formula	Molecular weight (g mol ⁻¹)	Water solubility (25°C) (mg L ⁻¹)
Fluometuron		C ₁₀ H ₁₁ F ₃ N ₂ O	232.2	90
Monolinuron		C ₉ H ₁₁ ClN ₂ O ₂	214.5	735
Diflubenzuron		C ₁₄ H ₉ ClF ₂ N ₂ O ₂	310.69	0.08

3. Development of the experimental setup, results and discussion

3.1. Development of the experimental setup

The two prototypes, AUTOPIF and AUTOPIF+ are developed, to track on-site the presence of non-native fluorescent pesticides in natural waters, by transforming it into PIF compounds after UV irradiation.

Sampling is done by a peristaltic pump and sent through a 25 mL cylindrical quartz photo reactor flow tube. It is irradiated on the way by a 125W Hg lamp (located at 5 cm of the tube), to create the PIF compounds. The irradiation time is control by adjusting the flow. The sample is then pumped through a cylindrical detection quartz flow cell (0.5 mL), and excited at 280 nm by a UV LED. This wavelength has been chosen because commercial UV LED lower than 280 nm are still not reliable nowadays for a long period of use. Although this irradiation

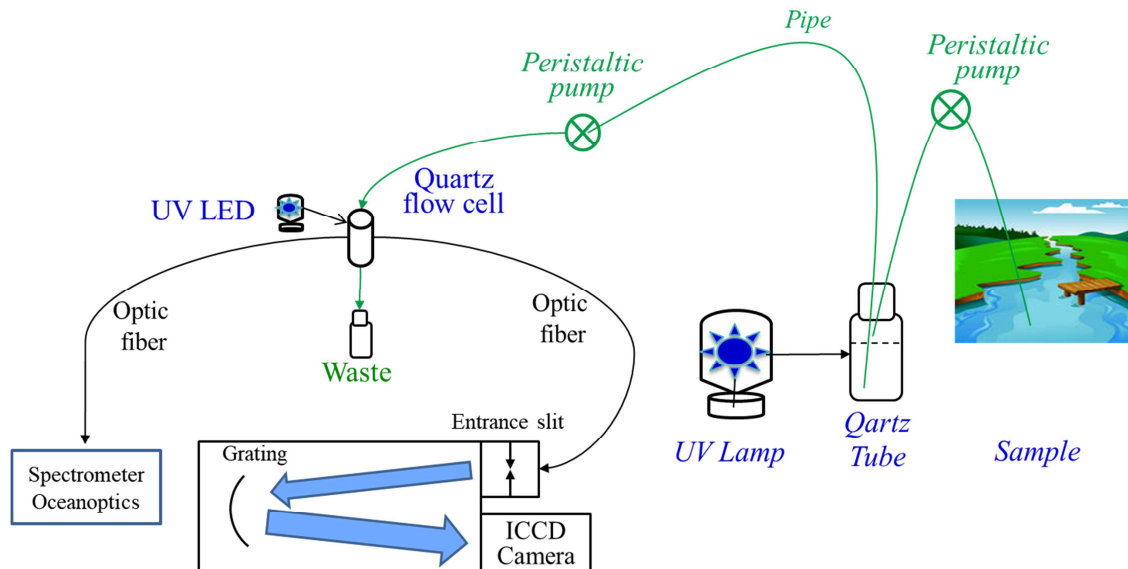
wavelength may not be the most appropriate for all pesticides studied but, it constitute a good compromise.

For AUTOPIF, the fluorescence is collected at 90° on one side by an optic fiber connected to a spectrometer (Ocean Optics) for the detection (Fig. 1). However, even if the spectrometer has a moderate cost, one of its drawbacks is due to its fixed sweeping zone between 200-400 nm. It will therefore be impossible to detect with the AUTOPIF system the spectra of fluorescents photoproducts with an emission wavelength greater than 400 nm.

For AUTOPIF+, to overcome the limitations of AUTOPIF, on the other side of the flow cell, a second optic fiber sends the fluorescence emission to a more resolute spectrometer (SpectraPro-550i), with a tunable sweeping zone from 200 to 800 nm (Fig. 1). Then the detection is done by an ICCD camera offering higher sensitivity. AUTOPIF+ is more resolute and more sensitive, but at a higher cost.

The two prototypes are design to be in the future integrated in an autonomous cabinet with a touch screen command interface.

Fig. 1. Description of the experimental systems “AUTOPIF” and “AUTOPIF+”



3.2 Photo-induced fluorescence properties

MLN, FLM and DFB have a low natural fluorescence while irradiation by UV rays resulted in the formation of highly fluorescent photoproducts. 20 mL of pesticides aqueous solutions ($1.5 \mu\text{g.mL}^{-1}$) have been pumped through the system.

One can see that in figure 2, MLN and DFB formed only one PIF compound at 360 nm and 410 nm respectively. FLM formed two different PIF compounds at 370 nm (PIF1) and at 430 nm (PIF 2). The photo-induced compounds are maximum after 1 min of irradiation for MLN and FLM-PIF1, after 2 min for DFB and after 3 min for FLM-PIF2 (Figure 3).

All these fluorescence parameters are sum-up in the Table 2. Same photoproduct was detected for DFB by Diaw et al ^[15] using classical-PIF method in mixture water-methanol (30:70, v/v) at pH4.

Fig 2. PIF emission spectra of MLN, FLM and DFB ($1.5 \mu\text{g.mL}^{-1}$) in aqueous medium **after one minute of irradiation** time and excitation wavelength at 280 nm.

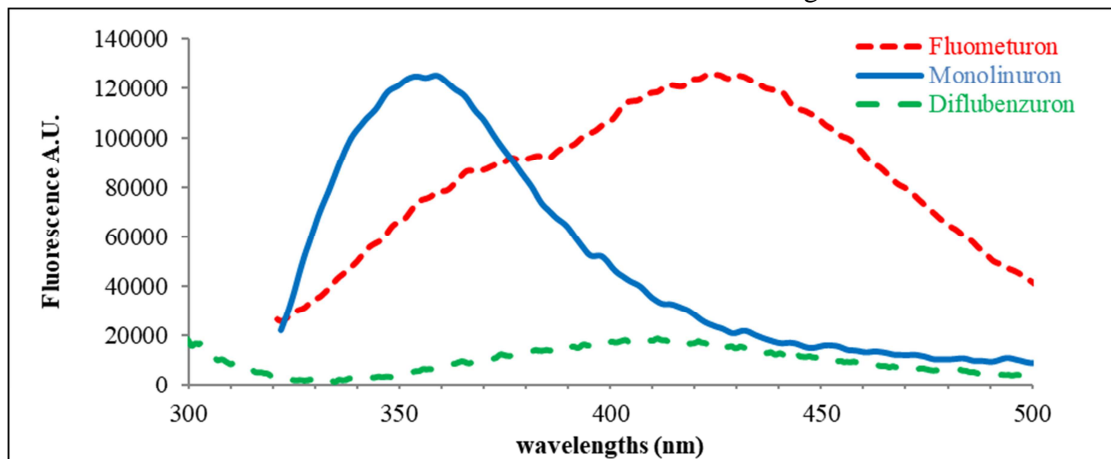


Fig 3. Formation of the photo-induced compounds as a function of the irradiation time.

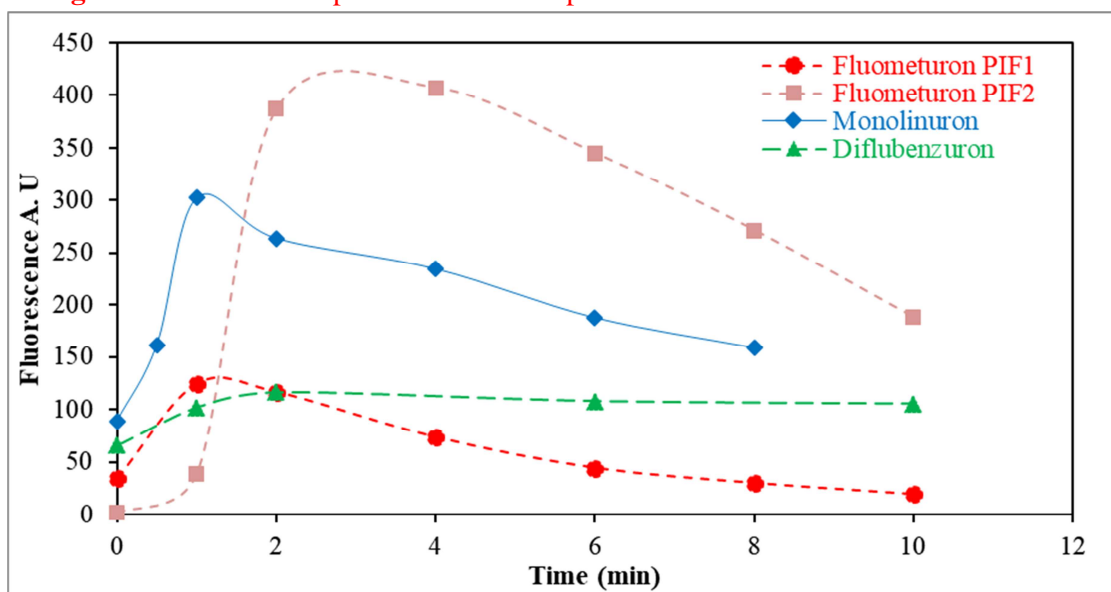


Table 2. Fluorescence spectral properties of the photoproducts of MLN, FLM and DFB ($1.5 \mu\text{g.mL}^{-1}$) in aqueous medium. **The more intense bands are in bold characters.**

Pesticides		λ_{ex} (nm)	λ_{em} (nm)
Monolinuron (MLN)	PIF 1	280	360
Diflubenzuron (DFB)	PIF 1	220/330	410
Fluometuron (FLM)	PIF 1	240	370
	PIF 2	230/310	430

3.3 Method Validation for pesticides determination

In order to evaluate the analytical usefulness of the method, analytical figures of merit were determined with AUTOPIF and AUTOPIF + (Figure 4). The study was performed with concentration ranges of $0.02\text{-}1.5 \mu\text{g mL}^{-1}$ for MLN, $0.02\text{-}2 \mu\text{g mL}^{-1}$ for FLM and $0.12\text{-}1 \mu\text{g mL}^{-1}$ for DFB. **The flow rate was adjusted to obtain an irradiation time of one min in the photo-reactor. This irradiation time was chosen as a compromise between the duration of the**

analysis and the sensitivity, although an irradiation time of 3 min could have given better sensitivity for FLM by studying its PIF2. Calibration curves were constructed by preparing samples in triplicate, containing increasing concentrations of each herbicide. For AUTOPIF, fluorescence intensity has been measured at 360 nm for MLN, and 370 nm for FLM-PIF1; as the spectrometer is limited to the wavelength range 200 to 400 nm, DFB cannot be studied. For AUTOPIF+, fluorescence intensity has been measured at 360 nm for MLN, 410 nm for DBF and for FLM we have recorded the emission of the PIF2 at 430 nm to obtain better sensitivity.

With AUTOPIF, the MLN give the most sensitive result with a slope of 232.16 AU. mL.mg⁻¹, while the FLM is less sensitive with a slope of 93.3 AU. mL.mg⁻¹. The calibration of DFB couldn't be realized with AUTOPIF because its photoproduct maximum emission wavelength is over 400 nm and therefore cannot be detected. With AUTOPIF+, the MLN give also the most sensitive result with a slope of 89116 AU. mL.mg⁻¹, while FLM and DFB are less sensitive, with a slope of 52832 AU. mL.mg⁻¹ and 16221 AU. mL.mg⁻¹ respectively.

The linearity of the calibration curve was evaluated by a variance analysis (table 3) for the three pesticides. The regression variance (V_{REG}) is in all cases significantly higher than the residual variance (V_{RES}) (p -value $\leq 5\%$), meaning that the regression is significant. Moreover, the lack of fit variance is not significantly higher than the pure error variance (p -value $> 5\%$) meaning that the linear model is validated. A Student t test has shown that the intercepts of calibration curves are not significantly different from zero (p -value $> 5\%$) for the three pesticides except the FLM for AQUAPIF+ which has p -value close to 3.3% (Table 4).

Measurements have been conducted using the AUTOPIF and AUTOPIF+ methods at the same time, on the same samples, to compare their results. With the AUTOPIF+ method we obtained detection limits 1.9, 5.2 and 17.4 ng.mL⁻¹ for MLN, FLM and DFB respectively which are lower with those obtained with the AUTOPIF method i.e.: 3.01 and 12.02 ng.mL⁻¹ for MLN and FLM; showing that AUTOPIF+ is more sensitive.

Moreover, the higher sensitivity obtains for MLN, for both AUTOPIF and AUTOPIF+, is also due to a better adequacy of the LED excitation wavelength (280 nm) to its optimal excitation wavelength (280 nm) compared to theses of DBF and FLM (i.e.: 220/330 nm; 240 nm; Table2).

Table 5 compares our results to some other on-site or laboratory methods for pesticide analysis, using UV detection or PIF. It sums up experimental protocols, analytical performances, advantages and drawbacks. We can note that AUTOPIF and AUTOPIF+ on-site systems give LOD values similar to the lowest of the literature obtained by laboratory systems without SPE pre-concentration; which indicates a good sensitivity. It should be noted that, compares to other method AUTOPIF and AUTOPIF+ are easy to implement, with a short analysis time and do not requires organic solvents.

Table 3: Evaluation parameters of the linear functions of the AUTOPIF and AUTOPIF+ method by variance analysis at a confidence level of 5%.

		AUTOPIF		AUTOPIF+		
		MLN	FLM	MLN	FLM	DFB
ANOVA 1	V_{REG}	$2.276 \cdot 10^5$	$6.481 \cdot 10^4$	$2.840 \cdot 10^{10}$	$2.242 \cdot 10^{10}$	$4.299 \cdot 10^8$
	V_{RES}	702.295	74.530	$2.748 \cdot 10^7$	$4.418 \cdot 10^6$	$2.806 \cdot 10^5$
	P value	0.0000	0.0000	0.0000	0.0000	0.0000
ANOVA 2	V_{LOF}	254.425	115.713	$2.477 \cdot 10^7$	$2.322 \cdot 10^6$	$13.552 \cdot 10^5$
	V_{PE}	836.657	62.175	$2.829 \cdot 10^7$	$5.180 \cdot 10^6$	$3.324 \cdot 10^5$
	P value	0.821	0.200	0.485	0.771	0.835

Table 4: Analytical figures of merit of the three pesticides and performances of AUTOPIF and AUTOPIF+ methods.

		AUTOPIF		AUTOPIF+		
		MLN	FLM	MLN	FLM	DFB
Slope		$2.32 \cdot 10^2$	93.30	$8.20 \cdot 10^4$	$5.33 \cdot 10^4$	$1.62 \cdot 10^4$
Intercept		1.85	7.13	$1.34 \cdot 10^3$	$1.85 \cdot 10^3$	$3.87 \cdot 10^2$
STD ^a		11.021	3.288	$2.18 \cdot 10^3$	$7.88 \cdot 10^2$	$2.43 \cdot 10^2$
P value		0.869	0.049	0.549	0.033	0.128
LOD (ng.mL ⁻¹) ^b		3.09	12.02	1.98	5.25	17.47
LOQ (ng.mL ⁻¹) ^c		10.29	37.08	8.49	16.75	55.42
LOD (mol.L ⁻¹) ^b		$14 \cdot 10^{-9}$	$46 \cdot 10^{-9}$	$8 \cdot 10^{-9}$	$22 \cdot 10^{-9}$	$56 \cdot 10^{-9}$
LOQ (mol.L ⁻¹) ^c		$51 \cdot 10^{-9}$	$155 \cdot 10^{-9}$	$25 \cdot 10^{-9}$	$69 \cdot 10^{-9}$	$169 \cdot 10^{-9}$

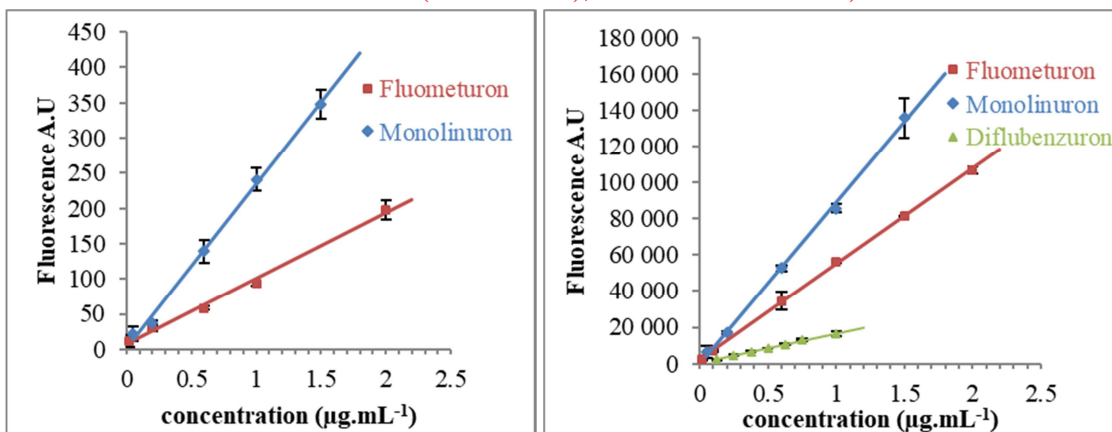
^a standard deviation of the intercept; ^b LOD: limit of detection, defined as the concentration of analyte giving a signal-to-noise (S/N) ratio of 3; ^c LOQ: limit of quantification, defined as the concentration of analyte giving a signal-to-noise(S/N) ratio of 10.

Table 5. Comparison of the analytical performances, protocols, advantages and drawbacks of some on-site systems and laboratory systems.

Pesticides	Method	Protocol	LOD (ng mL ⁻¹)	LQ (ng mL ⁻¹)	Sample	Advantages	Drawbacks	Ref.
MLN FLM DFB	On-site: “AUTOPIF” “AUTOPIF+”	PIF method implemented on a new automatic on-site detector in water without SPE pre-concentration.	1.9-12.0	5.9-36.0	- River water, - Tap water, - Sea water	- On-site system; - Continuous screening of pesticides. - Short analysis time. - Do not require organic solvents.		This work
Atrazine Isoproturon Diuron	On-site: “Aquapod SPE50”	SPE pre-concentration UV detection	0.9-1.5	-	River water	- On-site system. - Continuous screening of pesticides.	- Complex fluidic system - Need frequent maintenance.	[21]
Simazine Chloroluron Atrazine Isoproturon	On-site: “SAMOS”	SPE pre-concentration HPLC separation UV detection	0.6-1	-	River water	Selectivity	- Complex for an on-site system. - Need frequent maintenance.	[22]
Dichlorprop	Laboratory : PIF- Photooxydation	Photochemically induced fluorescence detection, correlated with the mechanism of photooxidation. 50 % (v/v) methanol and pH 5 buffer solutions	0.8	-	- Tomato; - Fruits (straw berry tree berry, orange, plum).	Sensitive	Longer irradiation time.	[26]
DFB	Laboratory : SPE-HPLC- PIF	On-line pre-concentration method. Fluorescence detection after photochemical induced fluorescence (PIF) postcolumn derivatization.	10	40	Ground water	Reduce the problem of interference	- Use of a lot of organic solvent; - Need complex pre-concentration steps.	[27]

Fenuron DFB	Laboratory : DL-PIF	Laser to obtain the photoproduct(s) and to simultaneously analyse their fluorescence. Short acquisition time on I CCD camera. Water and 50/50(v/v) methanol–water	1.5-4.8	5-16	- River water - Sea water	Sensitivity	Need a compromise between: - The laser wavelength, - The maximal absorption wavelength; - The optimal excitation wavelength.	[19]
Carboxin Monalide Propanil	Laboratory : HPLC-DL-PIF	On-line coupling of DL-PIF with liquid chromatography 60:40 (v:v) methanol-water.	0.53-3.64	1.77-12.14	- River water - Sea water	- Reduce the problem of interference; - Improve the selectivity	Need a compromise between: - The laser wavelength, - The maximal absorption wavelength; - The optimal excitation wavelength.	[28]
Isoproturon Néburon, Linuron, Diuron	Laboratory : FIA–MEPIF	Flow injection analysis. Micellar-enhanced photochemically induced fluorescence in buffered aqueous.	330-920	-	Tap water		Impossible to determine the maximum amount of photoinduced compounds	[29]
Isoproturon	Laboratory : SPE-HPLC-PIF	SPE pre-concentration. HPLC separation acetonitrile / buffer 60:40 (v / v) (pH 7, 0.01M). PIF derivatization. Fluorescence detection.	160-800	-	River water	Reduce interference problems	The time analysis is long	[7]
Lufenuron	Laboratory : PIF	Photo Induced Fluorescence. Performed in several media: methanol, ethanol and 2propanol	22	-	River water	Easy to implement	Time not defined in water and less sensitive than our method	[30]

Fig. 4. Calibration curves of MLN, FLM and DFB, (excitation at 280 nm UV LED, PIF fluorescence detection at 360 nm for MLN, 370 nm for FLM-PF1(AUTOPIF) and 430 nm for FLM-PIF2 (AUTOPIF+), and 410 nm for DFB).



3.4 Analytical applications

The usefulness of the method was tested with recovery studies carried out in different natural samples. From Senegal, we have taken samples from Senegal River which irrigates one of the most important agricultural areas of Senegal and can be submitted to some pesticide pollution; as river effluents can transfer the pesticide residue to sea water and induce fish contamination, studies have been also carried out on sea water samples from Dakar seaside. From France (Brittany district), we have taken samples from Penfeld River and from tap water of Brest city.

In order to get a matrix free of organic contaminants, the water samples were first filtered at 45 µm in order to eliminate the suspended organic matter, and then purified by solid phase extraction in order to eliminate the dissolved organic matter. The samples were then analysed by AUTOPIF and AUTOPIF+ using the standard addition method. They were initially fortified at 0.40 µg.mL⁻¹ for MLN and FLM and at 0.23 µg.mL⁻¹ for DFB. Increasing concentrations were added on those and the standard addition curves obtained were compared to the calibration curves (Figure 5 and 6).

Thus, in a first time, the linearity of the standard addition curves has been also tested by variance analysis (as explain in paragraph 3.3). In all cases, the regression is significant, without lack fit (at 5 % confidence level), meaning that the linear model is validated.

Then parallelism between the standard addition curves and the reference calibration curves was evaluated by a student *t* test. For each pesticide, the tabulated Student *t* value (confidence level of 5 %) is higher than the calculated *t* of the difference; showing **no significant differences between the slopes of the standard addition curves and the slope of the reference calibration curve** (Table 6 and 7 for AUTOPIF and AUTOPIF+ respectively). This indicates the absence of significant matrix effects.

Figure 5. Standard addition curves and the reference calibration of MLN.

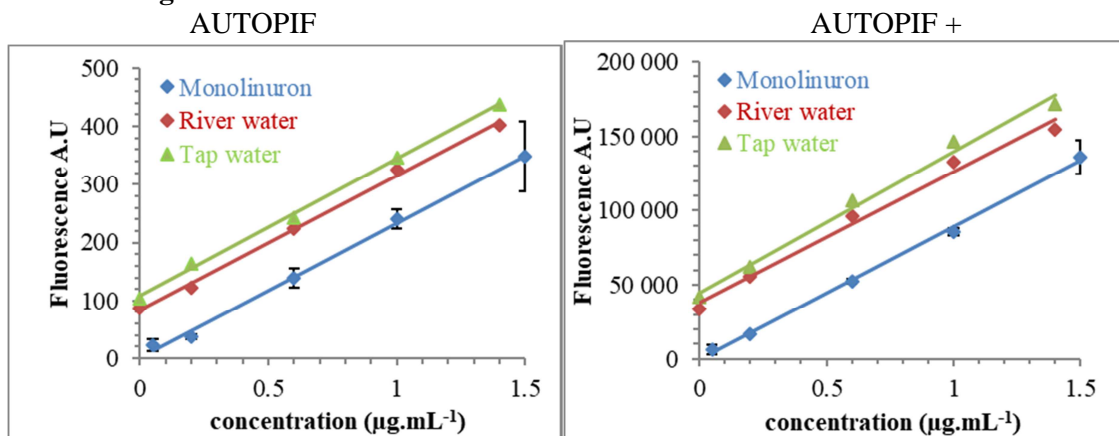


Figure 6. Standard addition curves and the reference calibration of FLM.

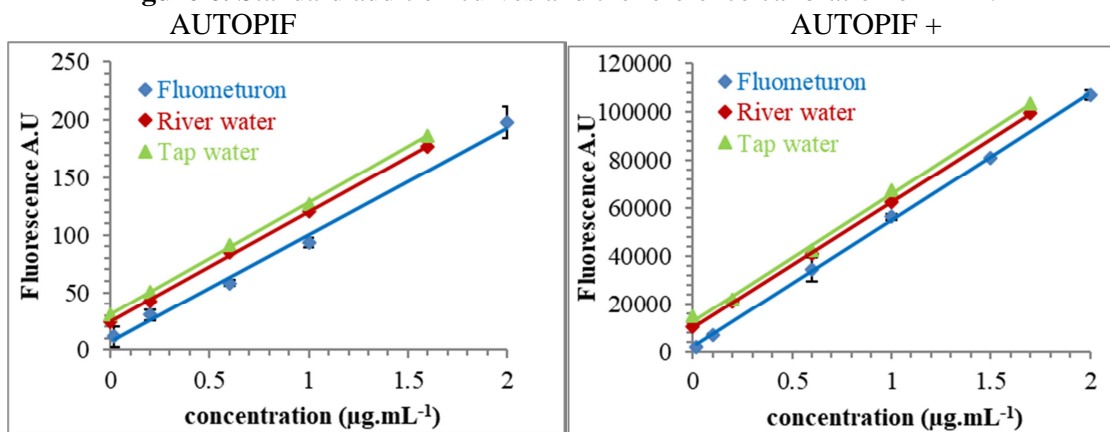


Table 6: Comparison of the addition standard curve and calibration curve slopes obtained by AUTOPIF method by a Student difference t test.

	AUTOPIF				AUTOPIF +	
	MLN	River water	Tap water		FLM	Tap water
Slope	232,16	236,05	231,34	:	98,52	96,44
STD	7,76	6,01	6,03	:	5,4	5,57
d.o.f		8	8	:		8
t_D		0,88621	0,186577	:		0,599521
t_S (5%)		2,306	2,306	:		2,306
SD		NO	NO	:		NO

STD: Standard Deviation of the slope of the standard addition curve; d.o.f.: degrees of freedom; t_D : Calculated Student value of the difference between the two slopes; t Tabulated Student t value; SD significant difference

Table 7: Comparison of the addition standard curve and calibration curve slopes obtained by AUTOPIF+ method by a Student difference t test.

	AUTOPIF+								
	MLN	River water	Tap water	FLM	River water	Tap water	DFB	Sea water	Tap water
Slope	$8.911 \cdot 10^4$	$8.787 \cdot 10^4$	$9.465 \cdot 10^4$	$5.266 \cdot 10^4$	$5.320 \cdot 10^4$	$5.176 \cdot 10^4$	$1.728 \cdot 10^4$	$1.766 \cdot 10^4$	$1.709 \cdot 10^4$
STD	$2.228 \cdot 10^3$	$5.803 \cdot 10^3$	$5.564 \cdot 10^3$	$1.130 \cdot 10^3$	$1.60 \cdot 10^3$	$2.084 \cdot 10^3$	$3.07 \cdot 10^2$	$5.44 \cdot 10^2$	$4.76 \cdot 10^2$
d.o.f		8	8		9	9		10	12
t_D		0.445	2.066		0.651	0.599		1.575	0.875
$t_S(5\%)$		2.306	2.306		2.262	2.262		2.28	2.179
SD		NO	NO		NO	NO		NO	NO

STD: Standard Deviation of the slope of the standard addition curve; d.o.f.: degrees of freedom; t_D : Calculated Student value of the difference between the two slopes; t Tabulated Student t value; SD significant difference

3.5 Method validation for applications

To validate the method a recovery study has been performed by spiking each water sample with an appropriate amount of MLN, FLM and DFB. The recoveries obtained can be observed in Table 8 for AUTOPIF and in Table 9 for AUTOPIF+. The recovery values were very close to 100%, ranging from 98% to 108 % for AUTOPIF and 99% to 105 % for AUTOPIF+. The relative standard deviation of the concentration found ranged from 3.5% to 7.6% for AUTOPIF and 2.3% to 5.1% for AUTOPIF+. The developed methods (AUTOPIF and AUTOPIF+) are then suitable to determine the benzoyl- and phenylurea pesticides in river, sea water and tap water samples, but the detection by AUTOPIF+ give the better results.

In order to show the practical interest of the method in these samples of sea and river water we have also done triplicates test points, at concentrations different from theses of the calibration curves. The predicted concentration values obtained by reference to the calibrations curves are found to be not different to the real value by a Student t test (Table 10 and 11).

Table 8: Recovery values obtained in spiked river water and tap water samples for the determination of MLN and FLM by AUTOPIF.

	Added ($\mu\text{g mL}^{-1}$)	Found ($\mu\text{g mL}^{-1}$)	Recovery, R (%)	Mean recovery, Rm (%)	Relative standard deviation, s(R) (%)
River water					
MLN	0.4	0.43 ± 0.08	107.50 ± 21.25		
	0.6	0.69 ± 0.08	115.00 ± 13.50		
	1.0	1.04 ± 0.09	104.00 ± 9.30	105.79	6.06
	1.4	1.38 ± 0.13	98.00 ± 9.29		
	1.8	1.87 ± 0.17	103.89 ± 9.44		
FLM	0.2	0.19 ± 0.11	95.00 ± 55.00		
	0.4	0.39 ± 0.15	97.50 ± 37.50		
	0.8	0.78 ± 0.13	97.50 ± 16.25	98.36	2.81
	1.2	1.19 ± 0.14	99.17 ± 11.67		
	1.9	1.95 ± 0.23	102.63 ± 12.11		
Tap water					
MLN	0.4	0.43 ± 0.08	107.50 ± 22.25		
	0.6	0.61 ± 0.08	101.67 ± 13.67		
	1.0	0.95 ± 0.08	95.00 ± 8.80	100.33	4.66
	1.4	1.38 ± 0.12	98.57 ± 8.57		
	1.8	1.78 ± 0.15	98.89 ± 98.89		
FLM	0.2	0.20 ± 0.10	100.00 ± 50.00		
	0.4	0.42 ± 0.16	105.00 ± 40.00		
	0.8	0.85 ± 0.15	106.25 ± 18.75	100.76	4.94
	1.2	1.18 ± 0.17	98.33 ± 14.17		
	1.9	1.79 ± 0.22	94.21 ± 11.58		

Table 9: Recovery values obtained in spiked river tap water and sea water samples for the determination of MLN, FLM and DFB “AUTOPIF+”

	Added ($\mu\text{g mL}^{-1}$)	Found ($\mu\text{g mL}^{-1}$)	Recovery, R (%)	Mean recovery, Rm (%)	Relative standard deviation, s(R) (%)
River water					
MLN	0.4	0.41 \pm 0.06	102.50 \pm 16.25		
	0.6	0.62 \pm 0.06	103.33 \pm 10.17		
	1.0	1.07 \pm 0.06	107.00 \pm 6.90	102.93	4.22
	1.4	1.48 \pm 0.08	105.71 \pm 5.93		
	1.8	1.73 \pm 0.11	96.11 \pm 6.11		
FLM	0.2	0.21 \pm 0.09	105.00 \pm 46.00		
	0.4	0.39 \pm 0.08	97.50 \pm 22.25	100.20	3.98
	0.8	0.76 \pm 0.07	95.00 \pm 9.50		
	1.2	1.23 \pm 0.07	102.50 \pm 6.00		
	1.9	1.92 \pm 0.11	101.05 \pm 5.79		
Sea water					
DFB	0.2	0.20 \pm 0.02	100.00 \pm 14.50		
	0.3	0.35 \pm 0.02	116.67 \pm 8.67		
	0.4	0.43 \pm 0.02	107.50 \pm 5.75	105.52	9.99
	0.5	0.56 \pm 0.02	112.00 \pm 5.20		
	0.7	0.64 \pm 0.02	91.43 \pm 3.86		
Tap water					
MLN	0.4	0.42 \pm 0.06	97.50 \pm 16.25		
	0.6	0.65 \pm 0.06	108.33 \pm 10.00		
	1.0	1.09 \pm 0.06	102.00 \pm 6.90	98.27	7.12
	1.4	1.38 \pm 0.08	90.71 \pm 5.93		
	1.8	1.91 \pm 0.11	92.78 \pm 6.11		
FLM	0.2	0.17 \pm 0.09	85.00 \pm 46.50		
	0.4	0.39 \pm 0.08	97.50 \pm 21.50		
	0.8	0.86 \pm 0.07	107.50 \pm 9.25	97.89	8.38
	1.2	1.23 \pm 0.07	102.50 \pm 6.00		
	1.9	1.84 \pm 0.08	96.84 \pm 4.53		
DFB	0.2	0.20 \pm 0.03	100.00 \pm 19.50		
	0.3	0.35 \pm 0.02	116.67 \pm 7.67		
	0.4	0.43 \pm 0.02	107.50 \pm 6.25	108.09	6.50
	0.5	0.56 \pm 0.02	112.00 \pm 4.60		
	0.7	0.73 \pm 0.02	104.29 \pm 3.57		

Table 10: Test points: Comparison of the concentration of each test point, to the concentrations found by regression, by a Student *t* test (AUTOPIF).

Pesticides	Sea water					River water				
	Added ($\mu\text{g mL}^{-1}$)	Found ($\mu\text{g mL}^{-1}$)	t_D	t_S (5%)	S D	Added ($\mu\text{g mL}^{-1}$)	Found ($\mu\text{g mL}^{-1}$)	t_D	t (5%)	SD
MLN	0.40	0.41 \pm 0.01	1.88	2.43	no	0.40	0.41 \pm 0.01	1.2 5	2.68	no
FLM	0.30	0.32 \pm 0.02	2.56	2.89	no	0.3	0.32 \pm 0.02	1.3 1	2.89	no

t_D : Calculated Student value of the difference between the two slopes; t Tabulated Student *t* value; SD significant difference

Table 11: Test points: Comparison of the concentration of each test point, to the concentrations found by regression, by a Student *t* test(AUTOPIF+).

Pesticides	Sea water					River water				
	Added ($\mu\text{g mL}^{-1}$)	Found ($\mu\text{g mL}^{-1}$)	t_D	$t_S(5\%)$	SD	Added ($\mu\text{g mL}^{-1}$)	Found ($\mu\text{g mL}^{-1}$)	t_D	$t(5\%)$	SD
MLN	0.40	0.41 \pm 0.02	1.772	2.300	no	0.40	0.41 \pm 0.02	1.309	2.230	no
FLM	0.30	0.32 \pm 0.02	2.566	2.716	no	0.30	0.32 \pm 0.02	1.520	2.743	no
DFB	0.30	0.31 \pm 0.05	0.779	2.320	no	0.30	0.31 \pm 0.05	0.251	2.352	no

t_D : Calculated Student value of the difference between the two slopes; t Tabulated Student *t* value; SD significant difference

4. Conclusion

In this present paper, we have associated, for the first time, the photo induced fluorescence method for the determination of pesticides in water, to new automatic **on-site alert systems** for environments waters, namely “AUTOPIF” and “AUTOPIF+”. **Thanks to the sensitivity of the fluorescence detection, we have reach the objective to develop a screening method while avoiding any pre-concentration step using complex fluidic systems as in some other commercial apparatus^[18, 19], in order to keep moderate cost.**

We have **use these prototypes** for the determination of benzoyl- and phenylurea pesticides i.e.: fluometuron, monolinuron and diflufenbuzon. **In future works, we will investigate in the identification of photoproducts responsible for the observed fluorescence.**

We obtained good analytical figures of merit, showing the ability of the methods for detection and quantification of these pesticides in the ng mL^{-1} range in aquatic environments **which is a sufficient value for an alert system.** The analytical applications in spiked natural waters of Senegal and Brest (France) have also led to satisfactory recovery values without significant matrix effects. AUTOPIF+ method gave better results with higher sensitivity thanks to the use of the ICCD camera detection. AUTOPIF is less sensitive but still at a sufficient level for an automatic alert system. In view of its analytical usefulness, AUTOPIF can be apply for the monitoring of pesticides in natural waters at a reasonable cost.

At this time, we are working on new evolutions of the prototypes while using other excitation wavelengths to increase sensitivity and specificity. We are also looking for an industrial partner in order to transform the prototypes into commercial systems and then to apply quality assurance protocol for its certification.

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